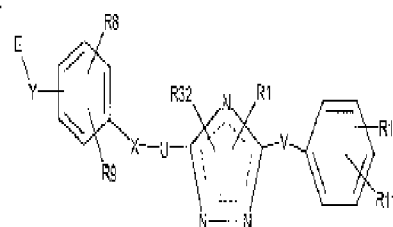


Serial No.: 10/580,202

Amendments to the Claims

What is claimed is:

1. (Canceled)
2. (Canceled)
3. (Currently Amended) A compound wherein the compound is of the Formula Ia:



Ia

and stereoisomers, or pharmaceutically acceptable salts, esters, and polymers thereof, wherein:

- (a) R1 is hydrogen;
- (b) R26, R27, R28 and R31 are each independently selected from the group consisting of hydrogen, hydroxy, cyano, amino, halo, oxo, C₁-C₃ alkyl, C₁-C₃ alkyl-COOR12, C₁-C₃ alkoxy, C₁ haloalkyl, C₁-C₃ haloalkoxy, C₁-C₃ cycloalkyl, aryl, aryl-C₁-C₃ alkyl, C(O)R13, COOR14, OC(O)R15, OS(O)R16, N(R17), NR18, C(O)R19, NR20, C(O)R21, SR22, S(O)R23, S(O)₂R24, and S(C₁-C₃ alkyl)R25); R12, R13, R14, R15, R16, R17, R18, R19, R20, R21, R22, R23, R24 and R25 are each independently selected from the group consisting of hydrogen, C₁-C₃ alkyl and aryl;
- (c) Y is selected from the group consisting of C₁-C₃ alkyl;
- (d) X is selected from the group consisting of a single bond;
- (e) U is an aliphatic linker wherein one carbon atom of the aliphatic linker is substituted with from one to two substituents each independently selected from R30;

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FILE COVERS 1907 - 30 Oct 2008 VOL 149 ISS 18
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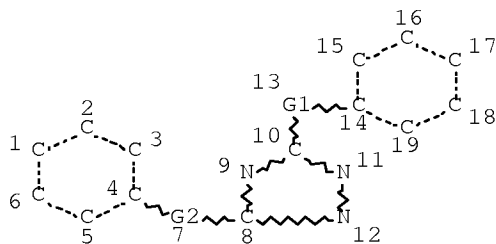
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L3 STR



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REP G2=(1-10) A

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DEFAULT ECLEVEL IS LIMITED

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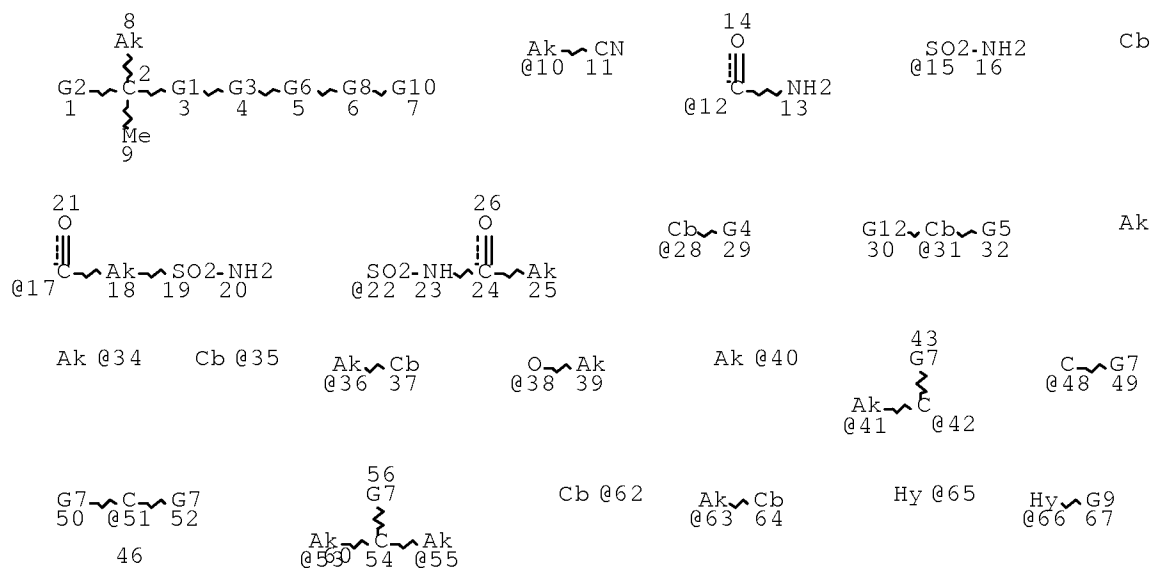
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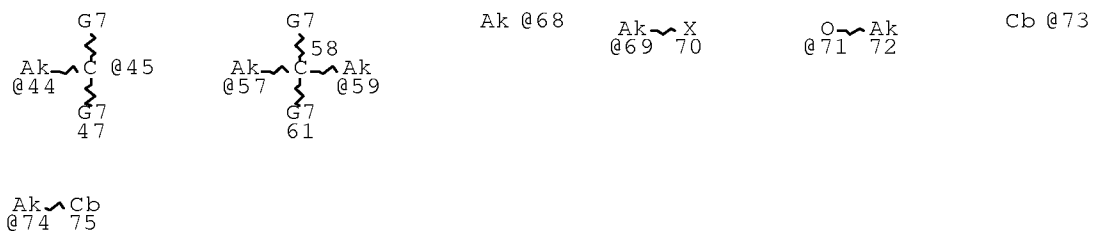


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@27

@33

Page 1-B



Page 2-A

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VAR G3=27/28/31

VAR G4=X/33/34/35/36/OH/38

VAR G5=34/35/36/OH/38

VAR G6=40/41-4 42-6/42-4 41-6/44-4 45-6/45-4 44-6/48/51/53-4 55-6/57-4 59-6

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VAR G9=X/68/69/71

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 reclassified documents, but they can be identified by 20060101/UPIC,
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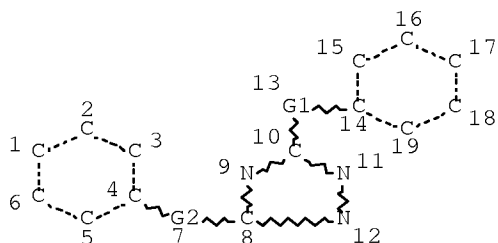
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>>> HELP for European Patent Classifications see HELP ECLA, HELP ICO <<<

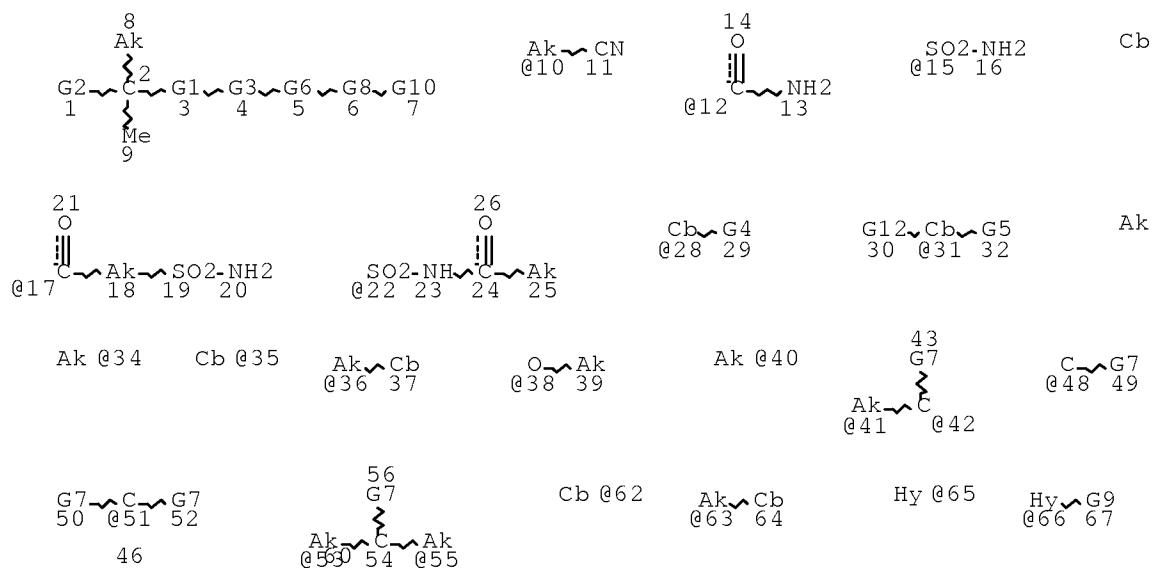
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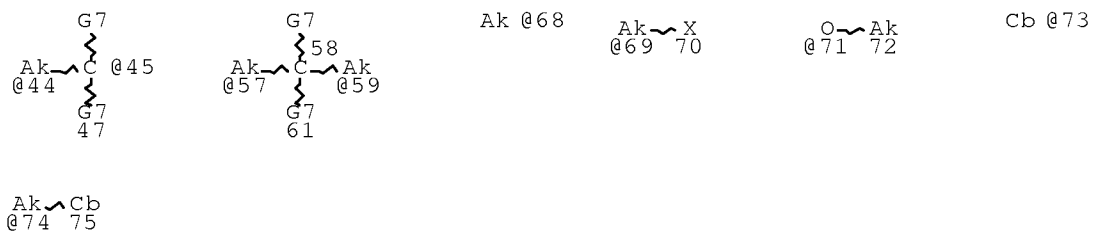


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@27

@33

Page 1-B



Page 2-A

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VAR G4=X/33/34/35/36/OH/38

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VAR G6=40/41-4 42-6/42-4 41-6/44-4 45-6/45-4 44-6/48/51/53-4 55-6/57-4 59-6

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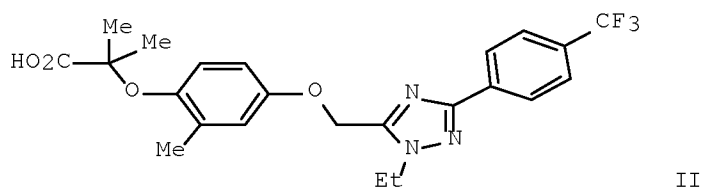
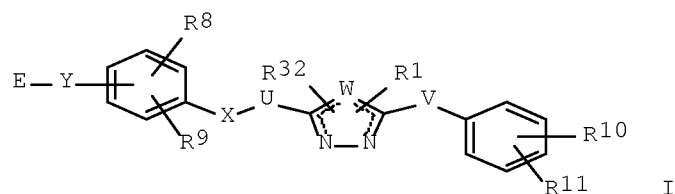
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 DOCUMENT NUMBER: 143:153383
 TITLE: Preparation of triazole, oxadiazole and thiadiazole
 derivatives as PPAR modulators for the treatment of

diabetes
 INVENTOR(S): Mantlo, Nathan Bryan; Navarro, Antonio; Saeed, Ashraf;
 Gernert, Douglas Linn; Ma, Tianwei; Pfeifer, Lance
 Allen
 PATENT ASSIGNEE(S): Eli Lilly and Company, USA
 SOURCE: PCT Int. Appl., 175 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005065683	A1	20050721	WO 2004-US39775	20041221
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EP 1725231	A1	20061129	EP 2004-812321	20041221
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			EP 2004-380158	A 20040721
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			EP 2004-350159	A 20040721
			WO 2004-US39775	W 20041221
OTHER SOURCE(S):	CASREACT 143:153383; MARPAT 143:153383			
GI				



AB The title compds. I [X = a single bond, O, S, SO₂ and N; U = an aliphatic linker; Y = O, C, S, NH and a single bond; W = N, O or S; E = CR₃R₄A or A (wherein A = carboxy, tetrazole, alkynitrile, carboxamide, sulfonamide and acylsufonamide; R₃ = H, alkyl, alkoxy; R₄ = H, alkyl, alkoxy, etc.; or R₃ and R₄ are optionally combined to form cycloalkyl); V = (hetero)alkyl, a bond; R₁ = H, alkyl, heteroaryl, etc.; R₈ = H, alkyl, alkenyl, halo; R₉ = H, alkyl, halo, etc.; R₁₀, R₁₁ = H, OH, CN, etc.; R₃₂ = a bond, H, halo, alkyl, etc.] which are modulators of peroxisome proliferator activated receptors (PPARs) and are useful for the treatment of diabetes and other metabolic disorders, were prepared and formulated. E.g., a multi-step synthesis of II, starting from Me glycolate and benzyl bromide, was given. The binding and cotransfection efficacy values for compds. I which are especially useful for modulating a PPAR receptor, are ≤ 100 nM and ≥ 50%, resp.

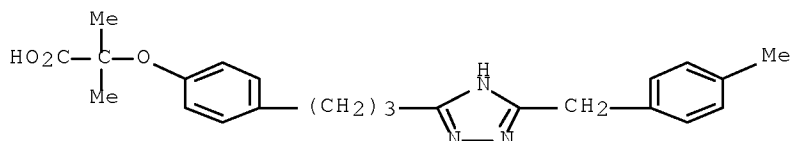
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860262-05-1P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of triazole, oxadiazole and thiadiazole derivs. as PPAR modulators for the treatment of diabetes)

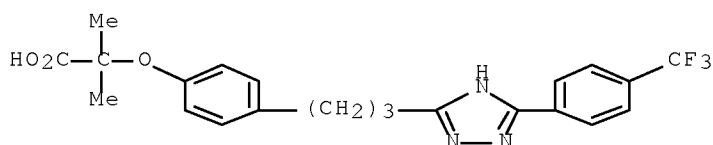
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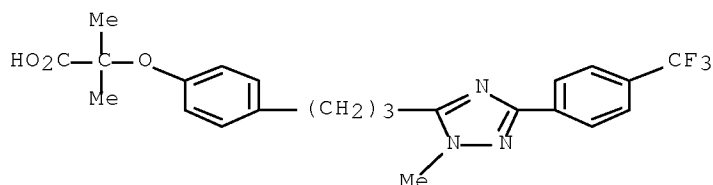
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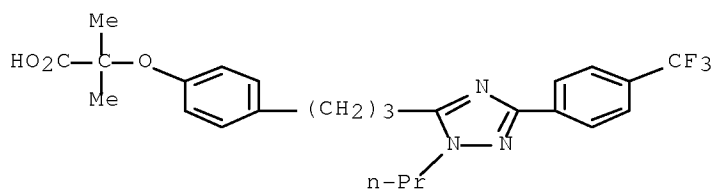
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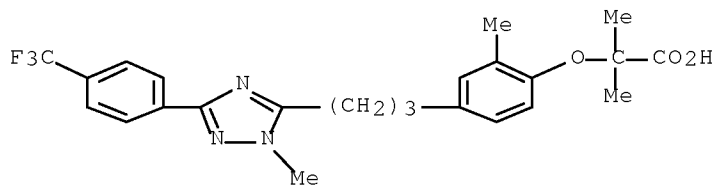
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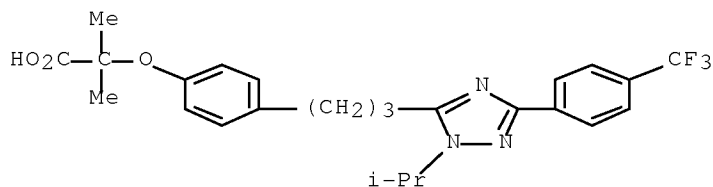
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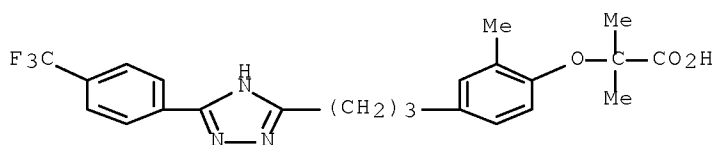
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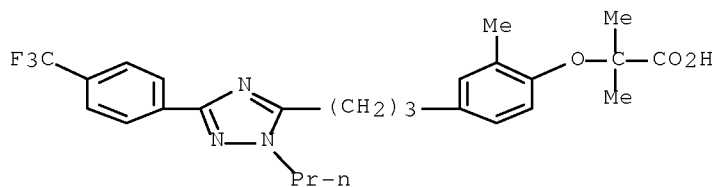
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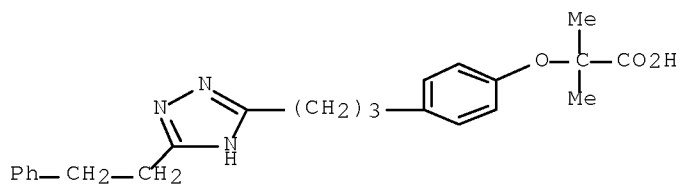
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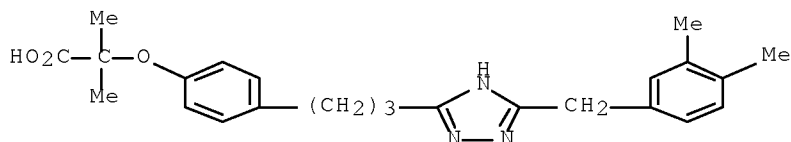


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RN 860262-05-1 CAPLUS
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